## **AMENDMENTS TO THE CLAIMS**

- 1. (Currently amended) A method of administering alprazolam or a pharmaceutically acceptable salt thereof to a mammal, comprising spraying the oral mucosa of the mammal with a propellant free buccal spray composition for transmucosal administration of alprazolam a pharmaceutically acceptable salt thereof comprising: alprazolam or a pharmaceutically acceptable salt thereof in an amount of between 0.001 and 60 percent by weight of the total composition; and a polar solvent in an amount between 30 and 99.69 percent by weight of the total composition.
- 2. (Currently amended) The composition method of claim 1, further comprising a taste mask and/or flavoring agent in an amount of between 0.1 and 10 percent by weight of the total composition.
- 3. (Currently amended) The composition method of claim 2, wherein the polar solvent is present in an amount between 37 and 98.58 percent by weight of the total composition, the alprazolam or a pharmaceutically acceptable salt thereof is present in an amount between 0.005 and 55 percent by weight of the total composition, and the taste mask and/or flavoring agent is present in an amount between 0.5 and 8 percent by weight of the total composition.
- 4. (Currently amended) The composition method of claim 3, wherein the polar solvent is present in an amount between 60.7 and 97.06 percent by weight of the total composition, the alprazolam or a pharmaceutically acceptable salt thereof is present in an amount between 0.01 and 40 percent by weight of the total composition, and the taste mask and/or flavoring agent is present in an amount between 0.75 and 7.5 percent by weight of the total composition.

Reply to Office Action of May 18, 2006

5. (Currently amended) The composition method of claim 1, wherein the polar solvent is selected from the group consisting of polyethylene glycols having a molecular weight between 400 and 1000, C<sub>2</sub> to C<sub>8</sub> mono- and poly-alcohols, and C<sub>7</sub> to C<sub>18</sub> alcohols of linear or branched configuration.

- 6. (Currently amended) The composition method of claim 1, wherein the polar solvent comprises polyethylene glycol.
- 7. (Currently amended) The <del>composition</del> method of claim 1, wherein the polar solvent comprises ethanol.
- 8. (Currently amended) The composition method of claim 2, wherein the flavoring agent is selected from the group consisting of synthetic or natural oil of peppermint, oil of spearmint, citrus oil, fruit flavors, sweeteners, and mixtures thereof.
  - 9. (Canceled).
- 10. (Currently amended) The method of claim [[9]]1, wherein the amount of the spray is predetermined.

Claims 11-21 (Canceled).

22. (Currently amended) A method of administering alprazolam or a pharmaceutically acceptable salt thereof to a mammal, comprising spraying the oral mucosa of the mammal with a propellant free buccal spray composition for transmucosal administration of alprazolam or a pharmaceutically acceptable salt thereof comprising: alprazolam or a pharmaceutically acceptable salt thereof in an amount between 0.005 and 55 percent by weight of the total composition; and a non-

polar solvent in an amount between 30 and 99<u>.69</u> percent by weight of the total composition.

- 23. (Currently amended) The composition method of claim 22, further comprising a taste mask and/or flavoring agent in an amount between 0.1 and 10 percent by weight of the total composition.
- 24. (Currently amended) The composition method of claim 23, wherein the flavoring agent is selected from the group consisting of synthetic or natural oil of peppermint, oil of spearmint, citrus oil, fruit flavors, sweeteners, and mixtures thereof.
- 25. (Currently amended) The composition method of claim 22, wherein the solvent is selected from the group consisting of (C<sub>2</sub>-C<sub>24</sub>) fatty acid (C<sub>2</sub>-C<sub>6</sub>) esters, C<sub>7</sub>-C<sub>18</sub> hydrocarbons of linear or branched configuration, C<sub>2</sub>-C<sub>6</sub> alkanoyl esters, and triglycerides of C<sub>2</sub>-C<sub>6</sub> carboxylic acids.
- 26. (Currently amended) The <del>composition</del> method of claim 25, wherein the solvent is a triglyceride.
  - 27. (Canceled).
- 28. (Currently amended) The method of claim [[27]]22, wherein the amount of the spray is predetermined.

Claims 29-39 (Canceled).

40. (Currently amended) A method of administering alprazolam or a pharmaceutically acceptable salt thereof to a mammal, comprising spraying the oral mucosa of the mammal with a buccal spray composition for transmucosal administration of alprazolam or a pharmaceutically acceptable salt thereof comprising:

alprazolam or a pharmaceutically acceptable salt thereof in an amount between 0.2 and 10 percent by weight of the total composition; and a polar solvent comprising propylene glycol and ethanol in an amount between 50 and 99.69 percent by weight of the total composition.

- 41. (Currently amended) A method of administering alprazolam or a pharmaceutically acceptable salt thereof to a mammal, comprising spraying the oral mucosa of the mammal with a propellant free buccal spray composition for transmucosal administration of alprazolam or a pharmaceutically acceptable salt thereof comprising: alprazolam or a pharmaceutically acceptable salt thereof in an amount of between 0.001 and 60 percent by weight of the total composition; and a mixture of a polar solvent and a non-polar solvent in an amount of between 30 and 99.69 percent by weight of the total composition, wherein the ratio of the polar solvent to the non-polar solvent ranges from 1:99 to 99:1.
- 42. (Currently amended) The composition method of claim 41, further comprising a taste mask and/or flavoring agent in an amount of between 0.1 and 10 percent by weight of the total composition.
- 43. (Currently amended) The composition method of claim 42, wherein the polar solvent is present in an amount between 37 and 98.58 percent by weight of the total composition, the alprazolam or a pharmaceutically acceptable salt thereof is present in an amount between 0.005 and 55 percent by weight of the total composition, and the taste mask and/or flavoring agent is present in an amount between 0.5 and 8 percent by weight of the total composition.
- 44. (Currently amended) The composition method of claim 43, wherein the polar solvent is present in an amount between 60.7 and 97.06 percent by weight of the

Reply to Office Action of May 18, 2006

total composition, the alprazolam or a pharmaceutically acceptable salt thereof is present in an amount between 0.01 and 40 percent by weight of the total composition, and the taste mask and/or flavoring agent is present in an amount between 0.75 and 7.5

percent by weight of the total composition.

45. (Currently amended) The composition method of claim 41, wherein the

polar solvent is selected from the group consisting of polyethylene glycols having a

molecular weight between 400 and 1000, C2 to C8 mono- and poly-alcohols, and C7 to

C<sub>18</sub> alcohols of linear or branched configuration and the non-polar solvent is selected

from the group consisting of (C2-C24) fatty acid (C2-C6) esters, C7-C18 hydrocarbons of

linear or branched configuration, C2-C6 alkanoyl esters, and triglycerides of C2-C6

carboxylic acids.

46. (Currently amended) The composition method of claim 42, wherein the

flavoring agent is selected from the group consisting of synthetic or natural oil of

peppermint, oil of spearmint, citrus oil, fruit flavors, sweeteners, and mixtures thereof.

47. (Canceled).

48. (Currently amended) The method of claim [[47]]41, wherein the amount of

the spray is predetermined.

Claims 49-56 (Canceled).

57. (Currently amended) [[A]]The method of claim 1, further comprising

treating anxiety in a patient, comprising spraying the oral mucosa of the patient with a

therapeutically effective amount of the buccal spray-of-claim 1.

6

Docket No.: N9810.0029/P029

- 58. (Currently amended) [[A]]The method of claim 11, further comprising treating anxiety in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 11.
- 59. (Currently amended) [[A]]<u>The</u> method of <u>claim 22</u>, <u>further comprising</u> treating anxiety in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of <u>claim 22</u>.
  - 60. (Canceled).
- 61. (Currently amended) [[A]]The method of claim 41, further comprising treating anxiety in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 41.
  - 62. (Canceled).
- 63. (Withdrawn and currently amended) [[A]]The method of <u>claim 1</u>, <u>further</u> <u>comprising</u> treating panic disorder in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of <u>claim 1</u>.
  - 64. (Canceled).
- 65. (Withdrawn and currently amended) [[A]]The method of claim 22, further comprising treating panic disorder in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of-claim 22.
  - 66. (Canceled).

Amendment dated November 20, 2006 Reply to Office Action of May 18, 2006

67. (Withdrawn and currently amended) [[A]]<u>The</u> method of <u>claim 41, further</u>

comprising treating panic disorder in a patient, comprising spraying the oral mucosa of

the patient with a therapeutically effective amount of the buccal spray-of claim 41.

68. (Canceled).

69. (Withdrawn and currently amended) [[A]]The method of claim 1, further

comprising inducing sleep in a patient, comprising spraying the oral mucosa of the

patient with a therapeutically effective amount of the buccal spray of claim 1.

70. (Canceled).

71. (Withdrawn and currently amended) [[A]]The method of claim 22, further

comprising inducing sleep in a patient, comprising spraying the oral mucosa of the

patient with a therapeutically effective amount of the buccal spray-of-claim 22.

72. (Canceled).

73. (Withdrawn and currently amended) [[A]]The method of claim 41, further

comprising inducing sleep in a patient, comprising spraying the oral mucosa of the

patient with a therapeutically effective amount of the buccal spray of claim 41.

74. (Canceled).

75. (Withdrawn and currently amended) [[A]]The method of claim 1, further

comprising treating the symptoms of premenstrual syndrome in a patient, comprising

spraying the oral mucosa of the patient with a therapeutically effective amount of the

buccal spray-of-claim-1.

76. (Canceled).

8

77. (Withdrawn and currently amended) [[A]]The method of claim 22, further comprising treating the symptoms of premenstrual syndrome in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of claim 22.

78. (Canceled).

79. (Withdrawn and currently amended) [[A]]The method of claim 41, further comprising treating the symptoms of premenstrual syndrome in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of-claim 41.

80. (Canceled).

81. (Withdrawn and currently amended) [[A]]The method of claim 1, further comprising treating chemotherapy induced emesis in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 1.

82. (Canceled).

83. (Withdrawn and currently amended) [[A]]The method of claim 22, further comprising treating chemotherapy induced emesis in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 22.

84. (Canceled).

85. (Withdrawn and currently amended) [[A]]<u>The</u> method of <u>claim 41</u>, <u>further</u> <u>comprising</u> treating chemotherapy induced emesis in a patient, comprising spraying

Docket No.: N9810.0029/P029

the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim-41.

86. (Canceled).

87. (Withdrawn and currently amended) [[A]]The method of claim 1, further comprising treating irritable-bowel syndrome in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 1.

88. (Canceled).

89. (Withdrawn and currently amended) [[A]]The method of claim 22, further comprising treating irritable-bowel syndrome in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray-of claim 22.

90. (Canceled)

91. (Withdrawn and currently amended) [[A]]The method of claim 41, further comprising treating irritable-bowel syndrome in a patient, comprising spraying the oral mucosa of the patient with a therapeutically effective amount of the buccal spray of claim 41.

92. (Canceled).